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PATENT COOPERATION TREATY

Translation

PCT Rec'd PCT/F10

8 JUN 2006

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

(Chapter II of the Patent Cooperation Treaty)

(PCT Article 36 and Rule 70)

Applicant's or a	gent's file referenc	e	-			
BIF023274/GP		FOR FURTHER A	ACTION	See Form PCT/IPEA/416		
International application No.		International filing d	ate (day/month/year)	Priority date (day/month/year)		
PCT/FR2004/000744		744 25.03.200)4	28.03.2003		
International Pa	International Patent Classification (IPC) or national classification and IPC					
Applicant INSTIT	UT PASTE	JR				
1. This under	report is the intern Article 35 and tra	national preliminary examination rensmitted to the applicant according	eport, established by this to Article 36.	International Preliminary Examining Authority		
2. This I	7					
3. This r	eport is also accon	npanied by ANNEXES, comprising		•		
a. [(sent to the	applicant and to the International B) Numanu) a tatal a f	shorter on Calling		
a. <u>L</u>				sheets, as follows: amended and are the basis for this report and/or		
	sheets Instruc	containing rectifications authorized	by this Authority (see Ru	ale 70.16 and Section 607 of the Administrative		
	sheets which supersede earlier sheets, but which this Authority considers contain an amendment that goes beyond the disclosure in the international application as filed, as indicated in item 4 of Box No. I and the Supplemental Box.					
Ъ. Г	(sent to the	International Bureau only) a total o	f (indicate type and numb	er of electronic carrier(c))		
_		mermanar Bar oan omy) a total o	(maleate type and number	or or electronic carrier(s))		
, containing a sequence listing and/or tables related thereto, in computer readable form only, as indicated in the Supplemental Box Relating to Sequence Listing (see Section 802 of the Administrative Instructions).						
4. This	report contains ind	ications relating to the following ite	ems:			
	Box No. I	Basis of the report				
	Box No. II	Priority				
	Box No. III	Non-establishment of opinion wi	th regard to novelty, inver	ntive step and industrial applicability		
	Box No. IV	Lack of unity of invention				
	Box No. V	Reasoned statement under Article citations and explanations support	e 35(2) with regard to nov ting such statement	elty, inventive step or industrial applicability;		
	Box No. VI	Certain documents cited				
	Box No. VII	Certain defects in the internation	al application			
	Box No. VIII	Certain observations on the inter-	national application			
Date of submission of the demand Date of completion of this report				his report		
				* · · · ·		
Name and mailing address of the IPEA/EP			Authorized officer			
Facsimile No.			Telephone No.			

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International application No.
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Box	No. I		Basis of the report		
1.	With	regard to	o the language, this report is based on the international er this item.	al application in the language in which it	was filed, unless otherwise
		which is	port is based on translations from the original language is the language of a translation furnished for the purpositernational search (Rule 12.3 and 23.1(b)) sublication of the international application (Rule 12.4) atternational preliminary examination (Rule 55.2 and/o	ses of:	·
2.	recei	regard to iving Office port): the intention the description in the de	to the elements of the international application, this refice in response to an invitation under Article 14 are transfer application as originally filed/furnished cription:	eport is based on (replacement sheets wh	tich have been furnished to the filed" and are not annexed to
		pages pages*	1-30	received by this Authority on	as originally filed/furnished
		pages*			
	\boxtimes	the clai			
		nos.	1-26		as originally filed/furnished
		nos.*			
		nos.*			
		nos.*			
	\boxtimes	the dra	wings:		
		sheets	1/7-7/7		as originally filed/furnished
		sheets*		received by this Authority on	
		sheets*		received by this Authority on	
	\boxtimes	a sequ	ence listing and/or any related table(s) - see Suppleme	ental Box Relating to Sequence Listing.	
3.		The ar	nendments have resulted in the cancellation of:		
			the description, pages		
			the claims, nos.		
			the drawings, sheets/figs		
		Ц	the sequence listing (specify):		····
			any table(s) related to sequence listing (specify):		
4.		This rethey h	eport has been established as if (some of) the amenda ave been considered to go beyond the disclosure as fil	ments annexed to this report and listed t ed, as indicated in the Supplemental Box	pelow had not been made, since (Rule 70.2(c)).
			the description, pages		
İ		$\overline{}$			
			the drawings, sheets/figs		
			the sequence listing (specify):		
	If it	em 4 apj	plies, some or all of those sheets may be marked "sup	erseded."	

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Box No. V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement				
1.	Statement			
	Novelty (N)	Claims	15-20, 26	YES
		Claims	13, 24	NO
	Inventive step (IS)	Claims	15-20, 26	YES
		Claims	1-14, 21-25	NO
	Industrial applicability	(IA) Claims	1-26	YES
		Claims		NO

2. Citations and explanations (Rule 70.7)

Reference is made to the following documents:

D1: WO 02/083892 A (MARLIERE PHILIPPE; POCHET SYLVIE (FR); BOUZON MADELEINE (FR); CENT) 24 October 2002 (2002-10-24);

- D2: WO 03/025163 A (KAMINSKI PIERRE-ALEXANDRE;

 MARLIERE PHILIPPE (FR); COTAYA RACHEL) 27 March
 2003 (2003-03-27);
- D3: DATABASE BIOSIS [online] BIOSCIENCES
 INFORMATION SERVICE, PHILADELPHIA, PA, US;
 2003, CHANG H K ET AL: "Directed evolution of
 Comamonas testosteroni GZ39 m-hydroxybenzoate
 hydroxylase for the synthesis of 4-substituted
 catechols." XP002297541 Database accession no.
 PREV200300546363;
- D4: WAN LIANGLU ET AL: "In vitro evolution of horse heart myoglobin to increase peroxidase activity" PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF THE UNITED STATES OF AMERICA, vol. 95, no. 22, 27 October 1998 (1998-10-27),

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pages 12825-12831, XP002297538 ISSN: 0027-8424.

D1 describes a method for evolving a protein X in such a way as to modify the properties thereof. More specifically, D1 makes reference to kinases belonging to classes EC 2.7.1, nucleotidyl transferases belonging to classes EC 2.7.7 and, in particular, polymerases and phosphorylases belonging to classes EC 2.4.2 (D1, claim 2).

The technical problem that document **D2** is intended to solve is that of producing nucleoside conversion enzymes and derivatives thereof that have <u>broader</u> enzymatic activity so as to enable diversification of the industrial production of these compounds (D2, page 2).

In order to provide recombinant enzymes capable of treating the widest variety of deviant substrates, recombinant N-deoxyribosyltransferases (EC 2.4.2.6) from various strains of *Lactobacillus* were isolated and cloned. This variety of enzyme has <u>broader specificity</u> and is produced via **mutations** in the wild-type genes or using chimeras of said wild-type genes (D2, page 3).

Moreover, it is already known that nucleoside analogues constitute a family of molecules that are active in the treatment of many bacterial, viral, parasitic and fungal infections as well as in anti-tumour chemotherapy (D2, page 1).

In view of documents D3 and D4, the use of directed in vitro evolution to enhance the properties of an enzyme or

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modify its specificity for a substrate are widely known in the prior art.

The present application does not fulfil the requirements set forth in PCT Article 33(1) because the subject matter of claims 13 and 14 does not comply with the requirement of novelty defined in PCT Article 33(2).

In light of the observations set out above, the subject matter of said claims cannot be differentiated from the disclosures in D1.

The present application does not fulfil the requirements set forth in PCT Article 33(1) because the subject matter of claims 1-12, 14, 21-23 and 25 does not involve an inventive step as defined in PCT Article 33(3).

The only difference between D1, which is considered to be the closest prior art, and claim 1 of the present international application is that, in D1, random mutagenesis is carried out *in vivo* in auxotrophic cells (D1, step (b)), while, in the present application, the DNA is mutated before being converted into auxotrophic cells.

The problem that the present invention is intended to solve can therefore be considered to be that of enhancing the prior art by providing an additional protein evolution method, in particular, for N-deoxyribosyltransferase (DTP).

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2.1 The solution proposed in claim 1 of the present application is not considered to be inventive (PCT Article 33(3)), for the following reasons:

In light of D2, the production of a DTP that has broadened enzymatic activity was an urgent requirement in view of the medical importance of nucleoside analogues.

It follows that, in light of D1, it would have been obvious for a person skilled in the art, aware of documents D3 and D4, to carry out the method in claim 1.

- 2.2 The same argument applies *mutatis mutandis* to the subject matter of corresponding independent claims 14 and 21, which are consequently not inventive either.
- 2.3 Dependent claims 2-12 do not contain any features which, in combination with the features of any one of the claims to which they refer, might define subject matter that fulfils the PCT requirement of inventive step (see documents D1 and D2 and the corresponding passages cited in the search report).
- 3. The subject matter of claim 26 is novel and the combination of features in claims 15-20 is not found in the prior art and cannot be derived in an obvious manner therefrom.

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	As a result, these claims fulfil the requirements		
	of PCT Article 33(2) and 33(3).		